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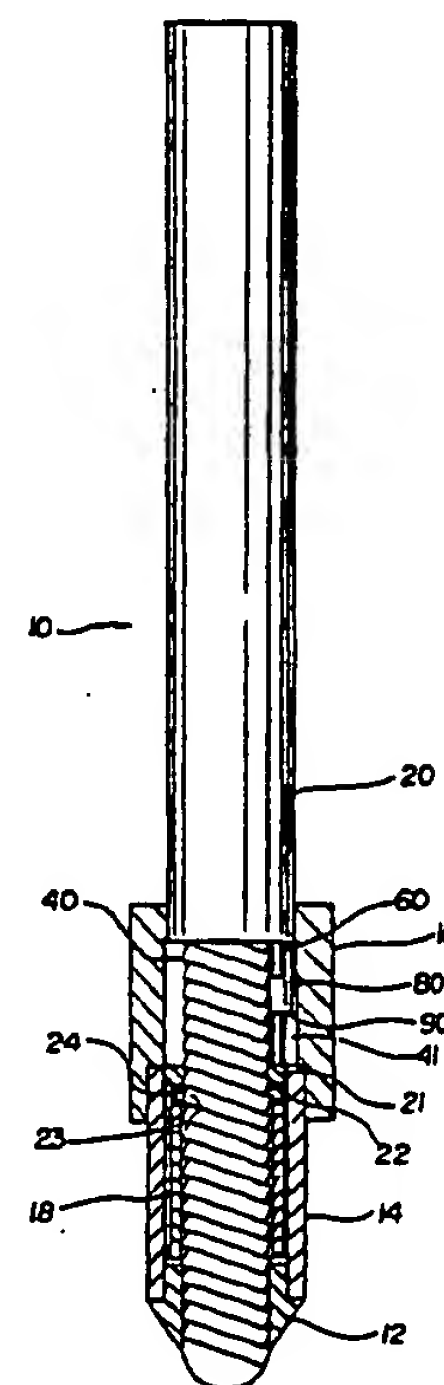
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61F 2/06	A2	(11) International Publication Number: WO 96/39998 (43) International Publication Date: 19 December 1996 (19.12.96)
<p>(21) International Application Number: PCT/US96/05842</p> <p>(22) International Filing Date: 26 April 1996 (26.04.96)</p> <p>(30) Priority Data: 08/484,006 7 June 1995 (07.06.95) US</p> <p>(71) Applicant: SCIMED LIFE SYSTEMS INC. [US/US]; One Scimed Place, Maple Grove, MN 55311-1566 (US).</p> <p>(72) Inventors: DEL TORO, Connie; 2165 Shenandoah Court, Unit H, Plymouth, MN 55447 (US). PETERSON, Dean, A.; 2904 17th Avenue North, Minneapolis, MN 55411 (US). SHOEMAKER, Susan, M.; 11106 190th Avenue N.W., Elk River, MN 55330 (US).</p> <p>(74) Agents: ANDERSON, William, E., II et al.; Vidas Arrett & Steinkraus, Suite 1540, 920 Second Avenue South, Minneapolis, MN 55402-4014 (US).</p>	<p>(81) Designated States: CA, JP, European patent (AT, BE, CH, DE, DK, ES, FL, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).</p> <p>Published <i>Without international search report and to be republished upon receipt of that report.</i></p>	

(54) Title: **PULL BACK SLEEVE SYSTEM WITH COMPRESSION RESISTANT INNER SHAFT**

(57) Abstract

A stent delivery system for delivering a stent comprising, a stent disposed on the distal end of the catheter, and an inner core, wherein the inner core is resistant to appreciable compression or accordion. The catheter further comprises a first sheath covering a portion of the inner core, wherein at least a portion of the distal end of the inner core is left uncovered by the first sheath, a retractable sheath which covers at least a portion of the stent and a portion of the distal end of the inner core and a retracting means for retracting the distal sheath to release the stent. The present invention also discloses a method of making the retractable sheath for a stent delivery system, comprising the steps of placing a sheath made of material comprising tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins on a mandrel, winding a wire coil around the Teflon® sheath, heating the tubing, cooling the tubing and removing the mandrel and coil.



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**PULL BACK SLEEVE SYSTEM WITH COMPRESSION
RESISTANT INNER SHAFT**

Field of the Invention

5 This invention relates to a stent delivery catheter system, such as the kind used in percutaneous transluminal coronary angioplasty (PTCA) procedures. More particularly, it relates to a stent delivery catheter employing a novel retractable protective sheath and a compression resistant inner shaft, and to a method of making the retractable protective sheath.

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Background of the Invention

 In typical PTCA procedures, a guiding catheter is percutaneously introduced into the cardiovascular system of a patient and advanced through the aorta until the distal end is in the ostium of the desired coronary artery. Using
15 fluoroscopy, a guide wire is then advanced through the guiding catheter and across the site to be treated in the coronary artery. An over the wire (OTW) balloon catheter is advanced over the guide wire to the treatment site. The balloon is then expanded to reopen the artery. The OTW catheter may have a guide wire lumen which is as long as the catheter or it may be a rapid exchange catheter wherein the
20 guide wire lumen is substantially shorter than the catheter. Alternatively, a fixed wire balloon catheter could be used. This device features a guide wire which is affixed to the catheter and cannot be removed.

 In certain known stent delivery catheters, a stent and an optional balloon are positioned at the distal end of the catheter, around a core lumen. The
25 stent and balloon are held down and covered by a sheath or sleeve. When the distal portion is in its desired location of the targeted vessel the sheath or sleeve is pulled back to expose the stent. After the sheath is removed, the stent is free to expand or be expanded. Such stent delivery catheters have had problems with the integrity of the inner core and the outer sheath. In a normal pull back system the friction
30 encountered when pulling the distal sheath off of the stent causes the innermost shaft to compress or accordion and the outermost sheath to elongate. This increases the likelihood of the inner core collapsing and the failure of the device to deploy the stent.

The present invention is directed toward remedying this collapsing or accordion type failure of the inner core. The invention is also directed toward an improved sheath and a method of making a low friction, strong, flexible sheath to be used in the stent delivery catheter.

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Summary of the Invention

The present invention provides an improved stent delivery catheter. The catheter includes a stent disposed on the distal end of the catheter, an inner core or inner lumen, which is flexible and resistant to appreciable compression or
10 accordion, and an outer sheath covering a majority of the inner core, excluding at least a portion of the distal end of the inner core. The catheter further comprises a retractable distal sheath which covers at least a portion of the stent and a portion of the distal end of the inner core and a retracting means for retracting the distal sheath to release the stent.

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The present invention further provides a retractable distal sheath and a method for making said retractable distal sheath. The inventive method comprises: placing a sheath comprising tetrafluoroethylene fluorocarbon polymers (TFEF) or fluorinated ethylene-propylene resins (FEP), such as Teflon™, on a mandrel, and winding a wire coil around the sheath. The sheath or tubing is then heated, allowing
20 the tubing to soften and the wire coil to create grooves in the soft tubing. After a certain period of heating, the tubing is allowed to cool and the mandrel and wire coil are removed. The resulting sheath demonstrates increased flexibility, sufficient strength and a low coefficient of friction.

Other objects, features, and characteristics of the present invention, as
25 well as the methods of operation and functions of the related elements of the structure, and the combination of parts and economics of manufacture, will become more apparent upon consideration of the following description with reference to the accompanying drawings, all of which form a part of this specification.

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Brief Description of the Figures

Figure 1 shows a side view of a catheter according to the invention including a cross-sectional view of the distal portion thereof.

Figure 2 shows a partial cut away view of a distal portion of a catheter according to the invention.

Figure 3 shows a side view of the proximal end of a catheter according to the invention showing the manifold portion thereof.

5 Figures 4a-4d show side views of optional contour patterns for the retractable distal sheath of the invention.

Detailed Description of the Invention

In Figure 1 there is shown a cross-section of the distal portion of a
10 specific embodiment of a stent delivery catheter generally designated as 10. The device generally comprises an outer sheath 20 which covers the majority of the catheter excluding a portion of the distal end of the catheter. This sheath 20 is characterized by a low friction coefficient and high flexibility, and preferably is comprised of a polyolefinic ionomer material, such as a single layer Surlyn™ sheath.
15 The outer sheath 20 surrounds an inner core 40 which extends to the distal tip 12 of the catheter. The inner core is preferably a spring coil 40, the manufacture of which is well known in the art, and is fashioned to be both flexible when navigated through body lumens and rigid when being pulled back upon itself during stent release. The spring coil may be made from a variety of material, including stainless
20 steel, elgiloy, Nitinol™, Kevlar™ or other metals and structural plastics. Preferably, it is made from stainless steel. The present invention further comprises a retractable distal sheath 14 covering a stent 18, which is loaded around the distal end of the inner core 40, and a retracting member 41, which is connected to the retractable distal sheath 14 and allows the physician to retract the distal sheath 14 from the
25 proximal end of the catheter. The retractable sheath 14 may be flexible or rigid, and is generally used to protect stent 18 and the vessel wall and/or to hold a self-expanding stent in the delivery configuration. The distal sheath 14 and the method for making it are discussed further below. The retracting member 41 may be a rod, a tube, a pull back wire or the like, but is preferably a wire. The retracting member
30 41 extends proximally through the outer sheath 20, preferably through a retracting member lumen 80, such as a tube preferably made from high density polyethylene (HDPE), but which could also be made from low density polyethylene (LDPE), polyimide, Teflon™ or other lubricious shaft material. In the preferred embodiment,

the retracting member lumen 80 extends longitudinally under the outer sheath 20, and houses the pull back wire 41. The retracting member lumen 80 that houses the pull back wire 41 may also carry flushing fluid for purging and cleaning the catheter at the distal end. Retracting member 41 exits the retracting member lumen 80 at exit hole 90, and continues distally to where it is attached to the distal sheath at point 21. The invention additionally comprises a proximal sheath 16 which covers the exposed area between the outer sheath 20 and the distal sheath 14, serving to protect the inner core 40 and the retracting member 10 in this area. The proximal sheath 16 is adhered to the proximal end of the distal sheath 14 and slidably overlaps the distal end of the outer sheath 20. As the distal sheath 14 is retracted, the proximal sheath 16 is forced back, sliding over the outer sheath 20 giving the distal sheath room to retract. The distance between the proximal end of the distal sheath 14 and exit hole 90 should preferably be far enough apart to allow complete release of the stent. The distal sheath 14 and the proximal sheath 16 may be two separate sheaths adhered to one another, or they may be combined to form one continuous sheath. Finally, a stiffening wire 60, preferably made from stainless steel, but which could also be made from Nitinol™ or Elgiloy, may also be incorporated longitudinally along the axis of the catheter 10 for extra stability and control.

Figure 2 shows the layers of the catheter excluding the distal portion of the outer sheath 20, the distal and proximal sheaths and the stent. As shown, the stiffening wire 60 and the retracting member lumen 80, which are positioned longitudinally along the catheter, may be truncated prior to the flexible distal tip. The truncated portion 28 may be terminated at the end of the outer sheath 20 or extend into the gap between the distal end of the outer sheath 20 and the proximal end of the distal sheath 14, as shown in Fig. 1. The retracting member 41 extends out through the truncated lumen 28 connecting with the distal sheath 14.

In the preferred embodiment, the distal sheath 14 is connected via a short section of hypotube 22, configured as an annular ring, to the pull back wire 41. The proximal end of the distal sheath 14 is adhered to the annular ring 22 and the pull back wire 10 is connected, preferably welded, to the inside of the annular ring 22. Proximal to the placement of the stent 18 is a stopper 24. The stopper 24 is usually a piece of tubing attached at position 23 to the inner core, and is used to

prevent the stent 18 from moving proximally when the distal sheath 14 is pulled back over the stent 18.

The proximal portion of the catheter, as shown in Figure 3, comprises of a manifold system 27 which includes a sliding member 26 slidably integrated
5 between the distal end of the manifold and the proximal Luer fitting 30. It is connected to the pull back wire 41 by a weld, insert mold or other connection means. By sliding the sliding member 26 of the manifold 27, distal to proximal, the distal sheath 14 is retracted exposing the stent 18. The manifold 27 may further
comprise a hydrating luer 32, which is located on the distal end of the manifold 27
10 and is used to hydrate the distal tip 12.

The inner core 40 is a non-compressible inner shaft that resists collapse or accordion type failure during the retraction of the distal sheath 14. In the preferred embodiment, a spring coil, most preferably a 6-fillar spring coil, is utilized for the inner core of the delivery device. A spring coil 14 such as used in
15 the present invention provides both flexibility during placement and rigidity during distal sheath retraction. The spring coil 14 allows the delivery system to deploy the stent 18 despite the amount of friction encountered at the distal end resulting from the use of a self expanding stent 18. As the wire 41 is pulled back to expose the self expanding stent 18, the spring coil 40 will collapse slightly upon itself until the
20 excess pitch has been taken up. Once this has happened, the spring coil 14 behaves as a rigid solid structure and therefore will not accordion, providing enough structural support for the distal sheath 14 to be pulled back and expose the self expanding stent 18.

To prepare the stent delivery catheter 10 the stent 18 is compressed
25 and loaded on the distal end of the inner core 40 inside of the distal sheath 14. The stent 18 is surrounded by protective distal sheath 14. The distal sheath remains covering the underlying stent during the placement of the stent 18 by the delivery catheter 10 through the patient's vasculature. During the placement of the stent, protective distal sheath 14 protects the patient's vasculature from the stent 18.
30 When it is time to expand the stent 18 into an enlarged diameter form and secure the stent in a patient's vasculature, the distal sheath 14 is retracted from over stent 18 by sliding the sliding member 26 proximally. As the sliding member is pulled back the distal sheath 14 starts to retract. Once the stent 18 is dragged slightly back by

the retracting distal sheath and is butted up against the stopper 24, the stent 18 expands fully as the distal sheath 14 continues to be pulled back. Preferably the stent is self expanding, such as a well known Nitinol™ stent, or it may be expanded by means of an optional internal balloon (not shown) positioned under the stent on the distal end of the inner core 40, as is well known in the art. Once the sheath 14 is fully retracted the optional placement balloon would be inflated through its inflation lumen (not shown) to deploy the stent. After the stent is expanded and in place, the catheter is withdrawn.

The stent deployment catheter preferably incorporates a distal sheath

10 material covering the stent with the following characteristics: low coefficient of friction to slide over the stent, which may comprise collagen material coating or bare metal, radial strength in order to hold down the self expanding stent and high flexibility to maneuver through torturous vasculature. Sheaths comprising tetrafluoroethylene fluorocarbon polymers (TFEF) or fluorinated ethylene-propylene
15 resins (FEP), such as Teflon™, have been found by the inventor to have the least amount of friction when dragged against the stent and inner core, while providing adequate radial strength to hold the stent in place. However, the TFEF/FEP sheaths have thick walls and make the distal tip too stiff for use in the peripheral anatomy. The present invention contemplates using TFEF/FEP sheaths as the distal sheath, or
20 both the distal sheath and the proximal sheath, of the stent deployment catheter and a new method of making the thick TFEF/FEP sheaths more flexible for use in a tortuous anatomy.

In making the desired distal sheath, a standard piece of Teflon™ tubing is placed on a mandrel just slightly smaller than the tubing's inner diameter. Using
25 a coil winder, a wire coil is wound directly over the tubing advancing from one end to the other end, noting the pitch and tension of the wire as the coil is laid on top of the Teflon™ tubing. The wire chosen can be either a round cross section or a rectangular cross section, preferably round with a diameter between .005" - .015". Heat is applied circumferentially to the coil wound tubing, at about 375 °C - 450
30 °C, preferably 420° C. The tubing is then allowed to cool to approximately room temperature and the spring coil and the mandrel are removed from the tubing leaving a contoured tube. The coil winder used to create the contoured surface may be wound to produce a variety of contour patterns. Figures 4a-4d illustrate possible

configurations. The amount of flexibility can be controlled by varying the amount of tension on the wire, the size of the wire, the wire profile, and the pitch of the wire. Preferably, a pitch of .010" - .075" is utilized.

5 This type of heat method provides a contoured surface on the Teflon™ sheath which results in a measurably improved retractable sheath having increased flexibility and sufficient strength. During heating, the tension from the hot wire leaves grooves in the softened tubing which allow the tubing to be more flexible. The resultant increase in flexibility is by approximately seven times when compared to the original piece of tubing, while still providing enough radial strength to hold
10 down the stent as well as providing the needed lubricity to remove the sheath from the stent. While fluorinated polymers are preferred, any thermoformable polymer may be employed.

The contouring process may also be used to provide flexible shafts for other medical devices such as balloon catheters or infusion catheters, or for any
15 other devices in which a flexible shaft is needed. In an infusion catheter, flexibility could be provided by contouring the distal end of the device. In a balloon catheter, the contouring could be used on either the inflation lumen or guidewire lumen as it would provide for fluid containment while providing flexibility.

The above disclosure is intended to be illustrative and not exhaustive.
20 These examples and description will suggest many variations and alternatives to one of ordinary skill in this art. All these alternatives and variations are intended to be included within the scope of the attached claims. Those familiar with the art may recognize other equivalents to the specific embodiments described herein which equivalents are also intended to be encompassed by the claims attached hereto.

What is claimed is as follows:

1. A stent delivery system comprising:
 - a substantially incompressible inner core having proximal and distal ends;
 - 5 a stent concentrically arranged around the inner core near the distal end;
 - a retractable sheath surrounding at least a portion of the stent, and retracting means connected to the distal sheath for retracting the distal sheath;
- 10 whereby when the retracting means is retracted the distal sheath is retracted freeing the stent for delivery.
2. The stent delivery system as in claim 1, wherein the inner core comprises a spring coil.
- 15 3. The stent delivery system as in claim 2, further comprising an outer sheath having proximal and distal ends and covering a portion of the inner core, wherein at least a portion of the distal end of the inner core is left exposed by the outer sheath, the retractable sheath overlapping a portion of the distal end of the outer sheath.
- 20 4. The stent delivery system as in claim 3, wherein the retracting means comprises a pull back wire attached to the retractable sheath, the pull back wire being longitudinally moveable.
- 25 5. The stent delivery system as in claim 4, further comprising an annular ring, wherein the pull back wire is attached to the annular ring and wherein the retractable sheath is adhered to the annular ring allowing the pull back wire to retract the retractable sheath.
- 30 6. The stent delivery system as in claim 5, further comprising a stopper positioned to prevent the stent from moving proximally as the retractable sheath is retracted to expose the stent.

7. The stent delivery system as in claim 6, further comprising a pull back wire lumen which partially encloses the pull back wire and is at least partially covered by the outer sheath.
- 5 8. The stent delivery system as in claim 7, further comprising a stiffening wire having proximal and distal ends positioned longitudinally along the inner core.
9. The stent delivery system as in claim 8, wherein the distal end of the stiffening wire and the distal end of the pull back wire lumen are truncated prior to
10 the annular ring.
10. The stent delivery system as in claim 5, wherein the retractable sheath comprises a proximal sheath having proximal and distal ends, the proximal end of the proximal sheath overlapping the distal end of the outer sheath, and a distal
15 sheath having proximal and distal ends, the distal end of the proximal sheath being connected to the proximal end of the distal sheath and wherein the annular ring is attached to the proximal end of the distal sheath.
11. The stent delivery system as in claim 10, further comprising a manifold, the
20 manifold comprising a sliding member which is connected to the pull back wire and acts to retract said pull back wire.
12. The stent delivery system as in claim 11, wherein the outer sheath comprises polyolefinic ionomer material.
25
13. The stent delivery system as in claim 12, wherein the retractable sheath is made of material which comprises tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins.
- 30 14. The stent delivery system as in claim 13, wherein the distal sheath comprises a contoured pattern of grooves.

15. The stent delivery system as in claim 14, further comprising a hydrating luer located on the manifold, a distal tip located at the distal end of the catheter and a communication means between the hydrating luer and the distal tip which allows for hydrating the distal tip from the hydrating luer.

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16. A retractable sheath for a stent delivery system comprising a tube of material made from tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins and which is contoured with a pattern of grooves, the contoured sheath being made from the process comprising the steps of:

- 10 placing the tube on a mandrel, wherein the diameter of the mandrel is slightly smaller than the inside diameter of the tube;
- winding a wire coil around the tube;
- heating the tube;
- cooling the tube; and
- 15 removing the mandrel and coil.

17. A method of making a retractable sheath for a stent delivery system, comprising the steps of:

- 20 placing a sheath made of material comprising tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins on a mandrel, wherein the diameter of the mandrel is slightly smaller than the inside diameter of the sheath;
- winding a wire coil around the sheath;
- heating the sheath;
- 25 cooling the sheath; and
- removing the mandrel and coil.

18. A method as in claim 17, wherein the wire coil is a cross-section wire coil.

30 19. A method as in claim 18, wherein the wire coil is a rectangular wire coil.

20. A stent delivery system comprising:

a substantially incompressible inner core having proximal and distal ends, and being made from a wire coil with a predetermined pitch such that it is flexible in an uncompressed state, and will become rigid when compressed;

5 a stent concentrically arranged around the inner core near the distal end;

a retractable sheath surrounding at least a portion of the stent, and retracting means connected to the distal sheath for retracting the distal sheath;

10 whereby when the retracting means is retracted to free the stent for delivery, the distal sheath is retracted, causing the wire coil to compress and taking up the excess pitch in the wire coil, making the wire coil substantially rigid, preventing the inner core from collapsing.

Fig. 1

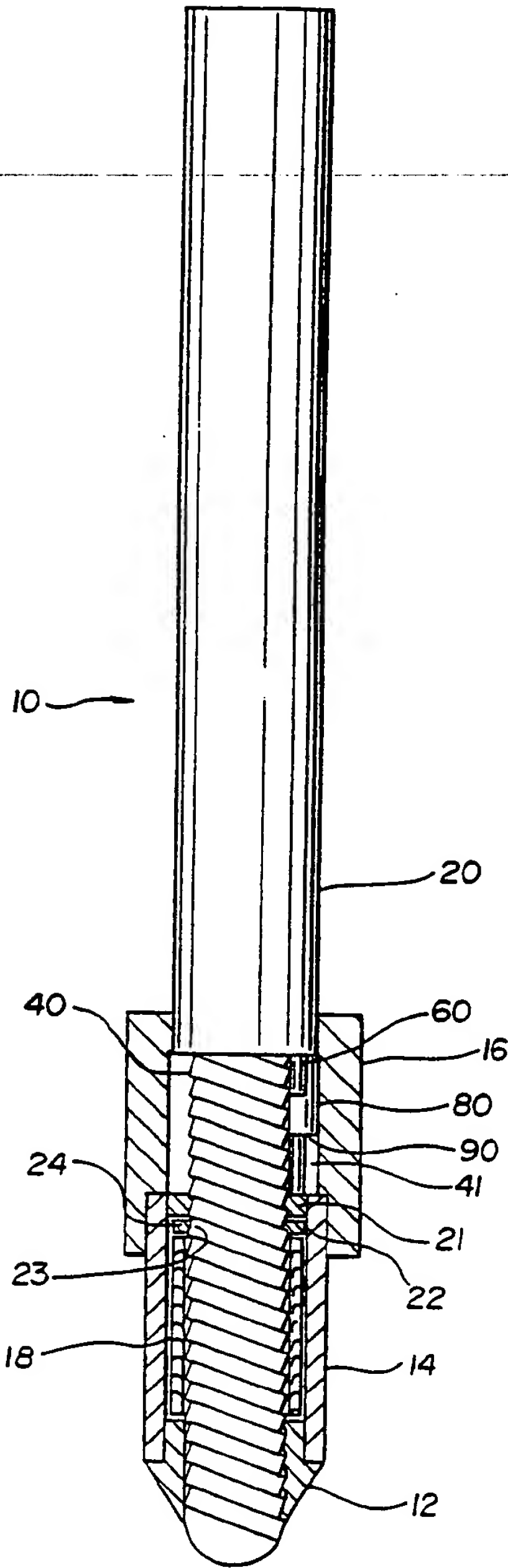
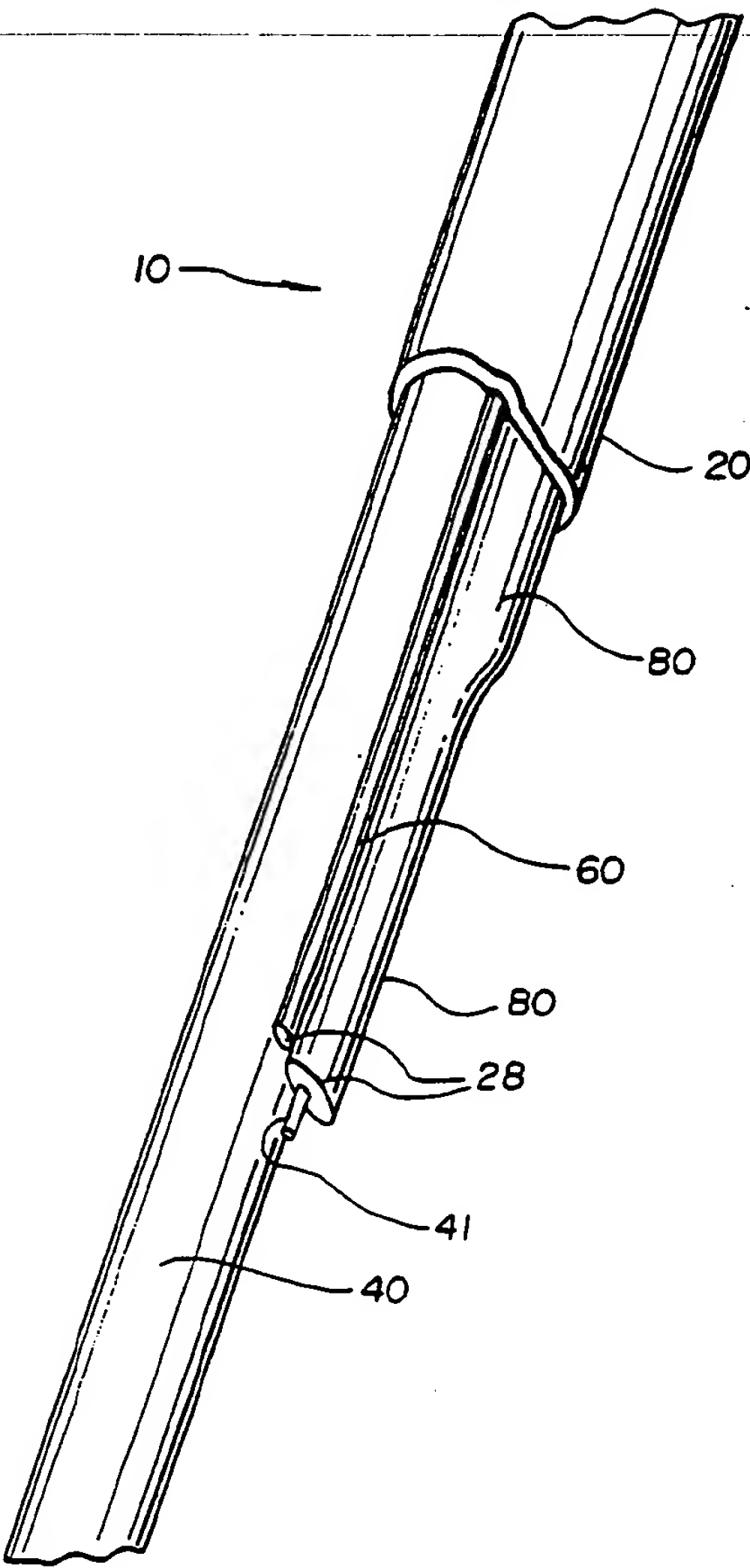
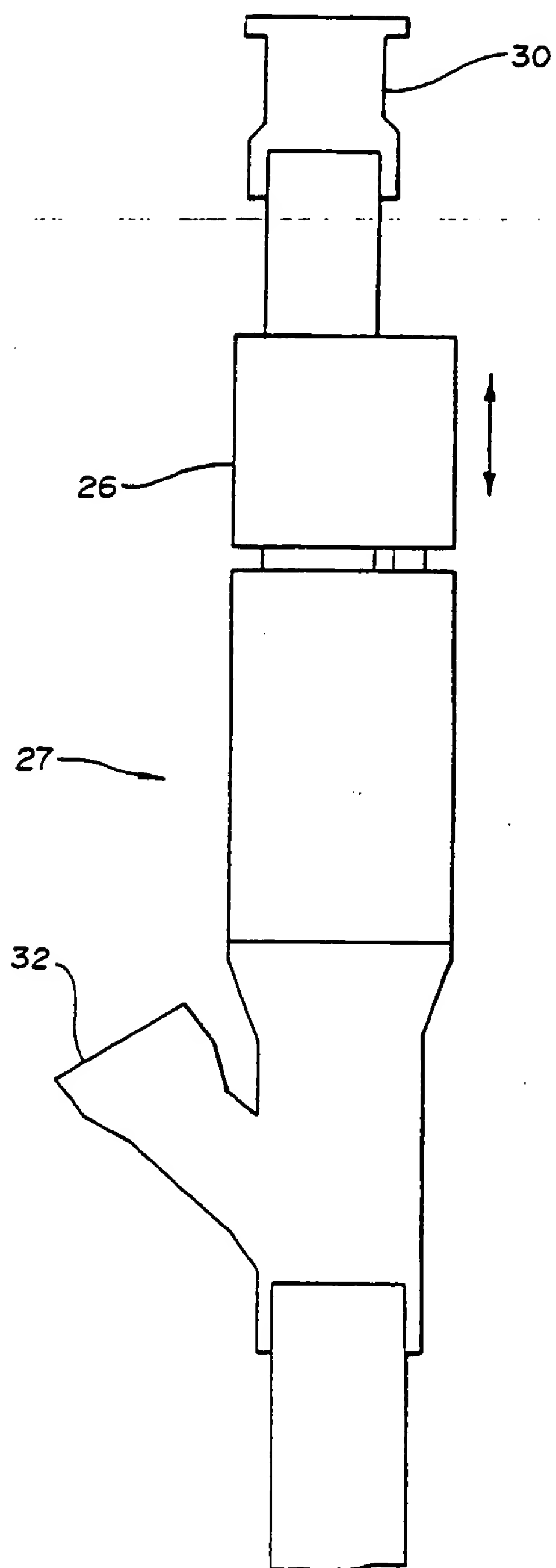


Fig. 2



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Fig. 3

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Fig. 4a

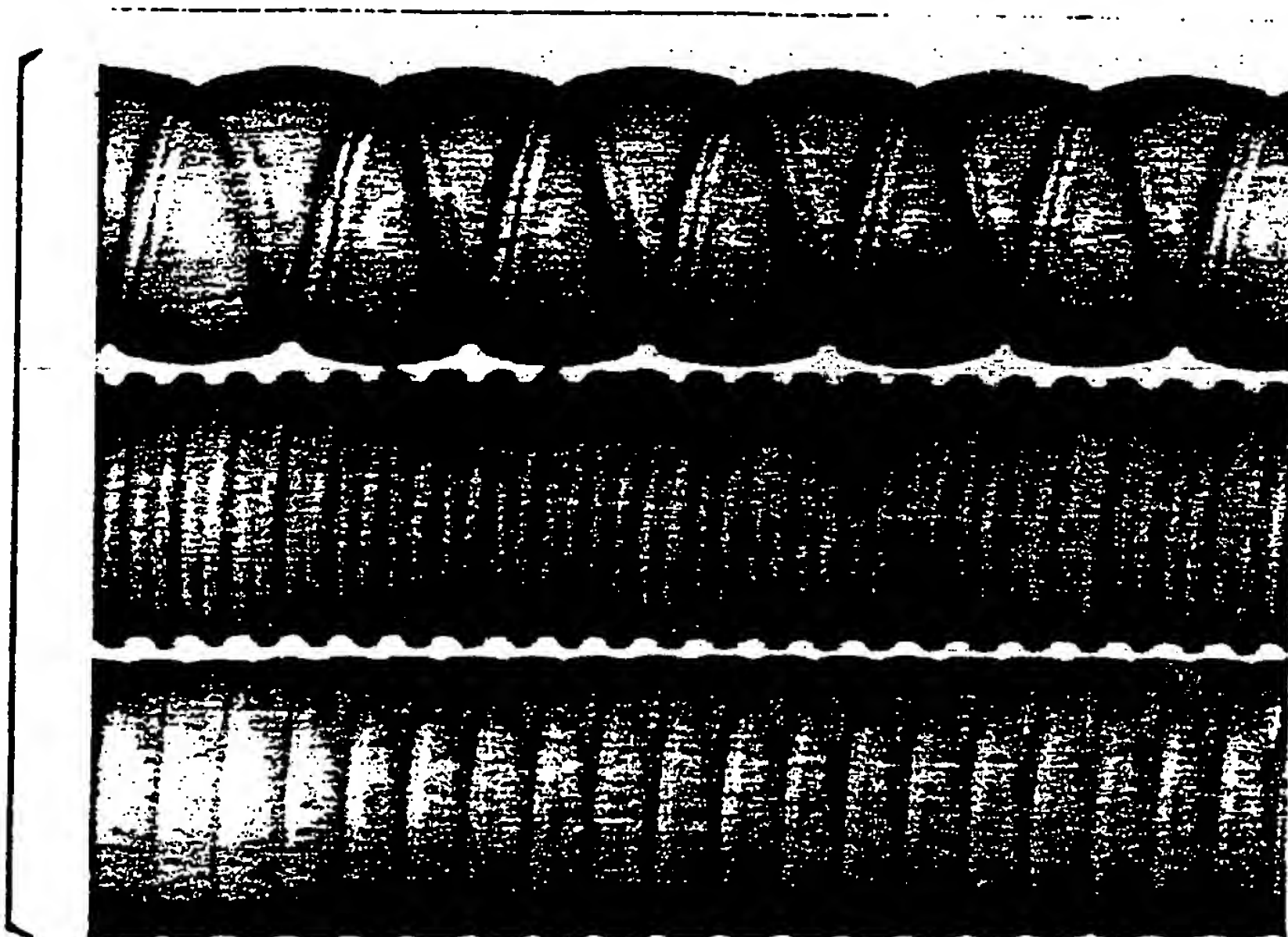
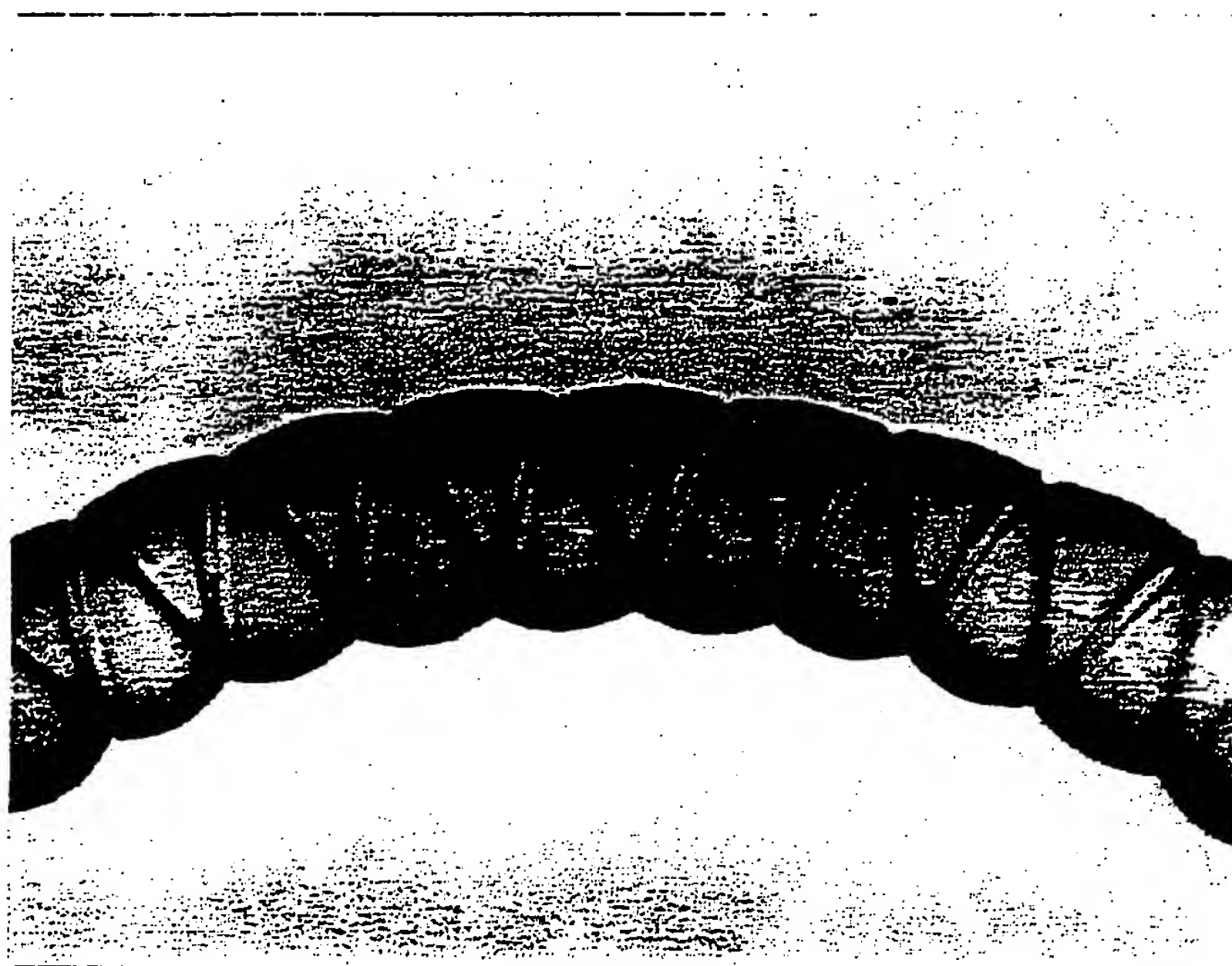


Fig. 4b



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Fig. 4c

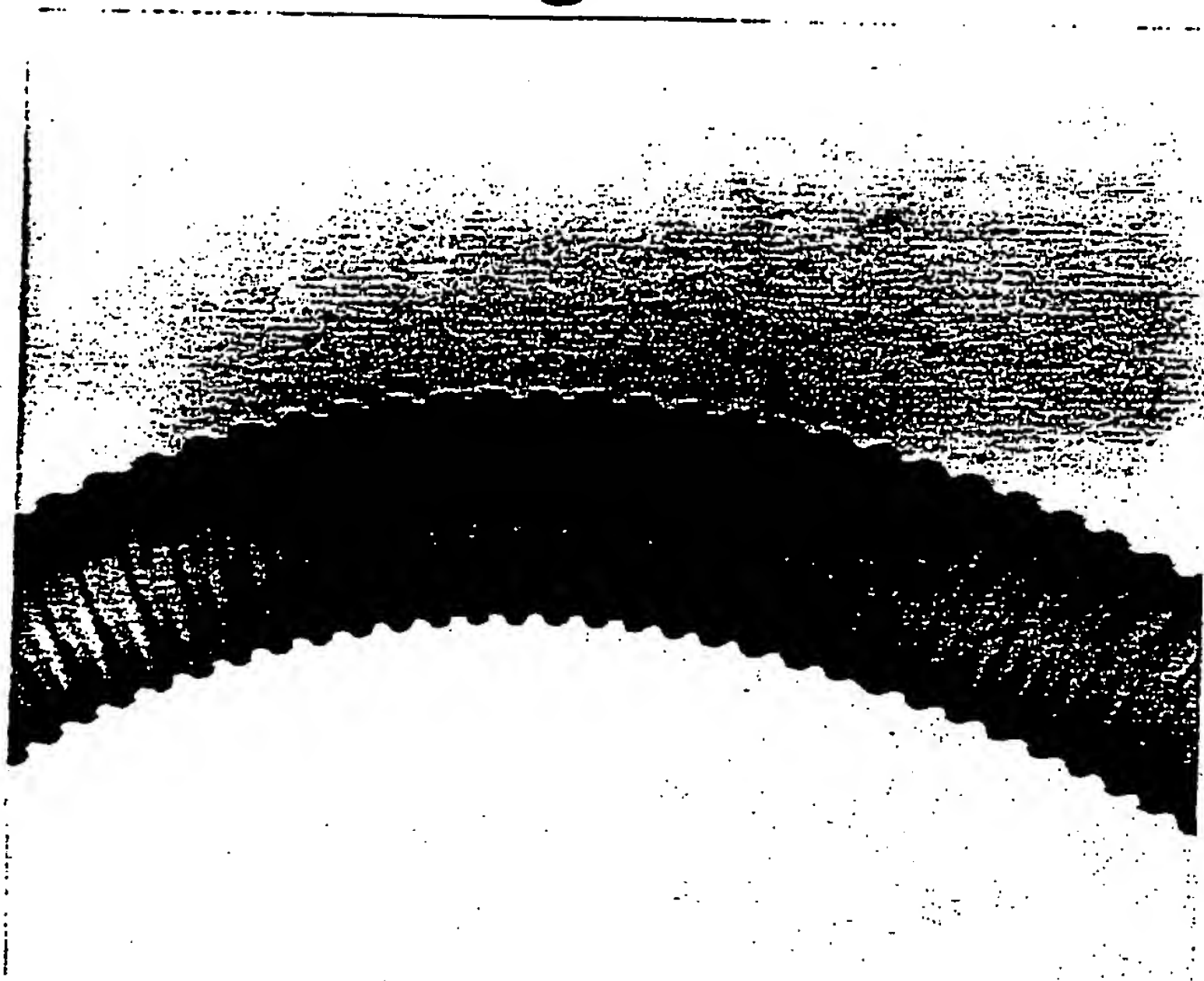
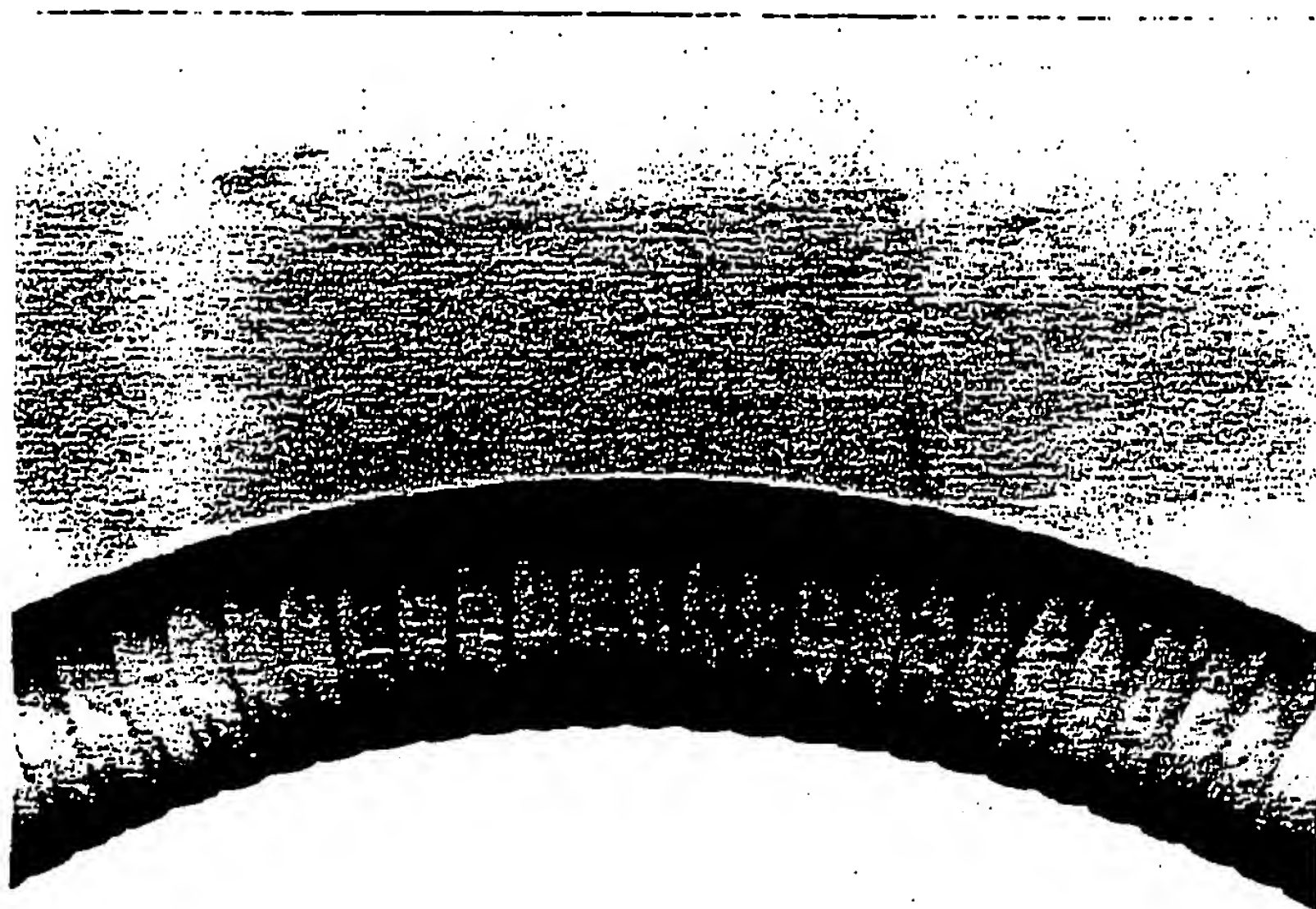


Fig. 4d



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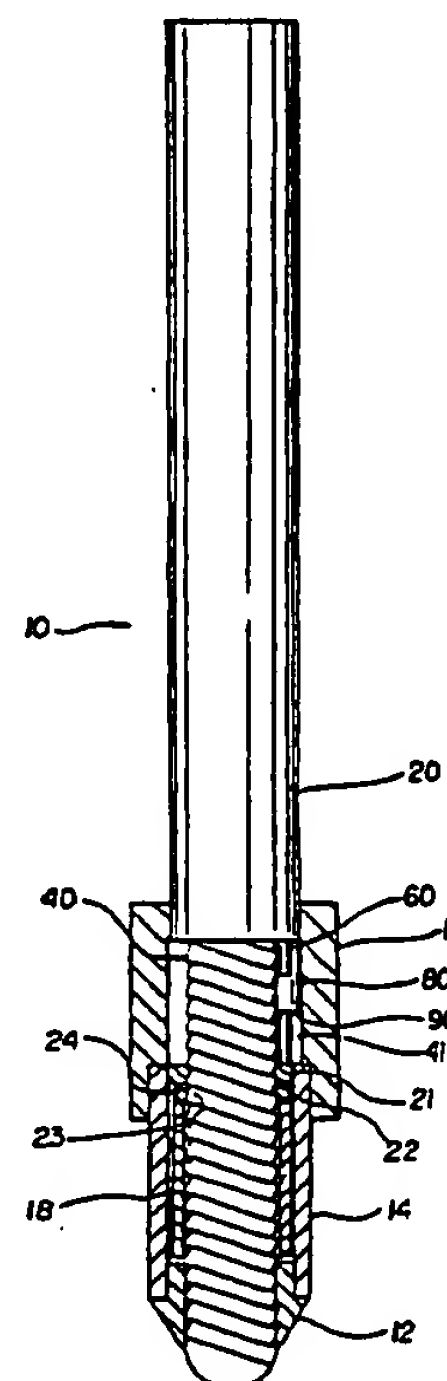
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(51) International Patent Classification ⁶ : A61F 2/06		A3	(11) International Publication Number: WO 96/39998
			(43) International Publication Date: 19 December 1996 (19.12.96)
(21) International Application Number: PCT/US96/05842		(81) Designated States: CA, JP, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(22) International Filing Date: 26 April 1996 (26.04.96)			
(30) Priority Data: 08/484,006 7 June 1995 (07.06.95) US		Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.	
(71) Applicant: SCIMED LIFE SYSTEMS INC. [US/US]; One Scimed Place, Maple Grove, MN 55311-1566 (US);		(88) Date of publication of the international search report: 23 January 1997 (23.01.97)	
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(54) Title: PULL BACK SLEEVE SYSTEM WITH COMPRESSION RESISTANT INNER SHAFT

(57) Abstract

A stent delivery system for delivering a stent comprising, a stent disposed on the distal end of the catheter, and an inner core, wherein the inner core is resistant to appreciable compression or accordion. The catheter further comprises a first sheath covering a portion of the inner core, wherein at least a portion of the distal end of the inner core is left uncovered by the first sheath, a retractable sheath which covers at least a portion of the stent and a portion of the distal end of the inner core and a retracting means for retracting the distal sheath to release the stent. The present invention also discloses a method of making the retractable sheath for a stent delivery system, comprising the steps of placing a sheath made of material comprising tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins on a mandrel, winding a wire coil around the Teflon® sheath, heating the tubing, cooling the tubing and removing the mandrel and coil.



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INTERNATIONAL SEARCH REPORT

Inten. Application No
PCT/US 96/05842

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61F2/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	EP 0 556 940 A (FISCHELL) 25 August 1993 see column 4, line 4 - column 5, line 28; figures	1 20
A	WO 93 11823 A (TARGET THERAPEUTICS) 24 June 1993 see abstract; figures	1,2,20
A	US 4 795 458 A (REGAN) 3 January 1989 see column 3, line 46 - column 4, line 23; figures	1,2,20
E	WO 96 13228 A (SCHNEIDER) 9 May 1996 see abstract; figures 4-8	1,2

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

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- *O* document referring to an oral disclosure, use, exhibition or other means
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- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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Date of the actual completion of the international search

30 September 1996

Date of mailing of the international search report

12.12.96

Name and mailing address of the ISA

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KOUSOURETAS, I

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 96/05842

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. Claims: 1-15,20 A stent delivery system comprising an incompressible inner core, a stent, a retractable sheath and retracting means.
2. Claims: 16-19 A retractable sheath made from tetrafluorethylene fluorocarbon polymers, which is contoured with a pattern of grooves, the contoured sheath being made from a described process.

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-15,20

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 96/05842

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A-556940	25-08-93	DE-A- 3786721	02-09-93
		EP-A- 0257091	02-03-88
		WO-A- 8704935	27-08-87
		US-A- 4768507	06-09-88

WO-A-9311823	24-06-93	US-A- 5234437	10-08-93
		AU-A- 3238293	19-07-93
		CA-A- 2125652	24-06-93
		EP-A- 0616544	28-09-94
		JP-T- 7503153	06-04-95

US-A-4795458	03-01-89	EP-A- 0362444	11-04-90

WO-A-9613228	09-05-96	NONE	

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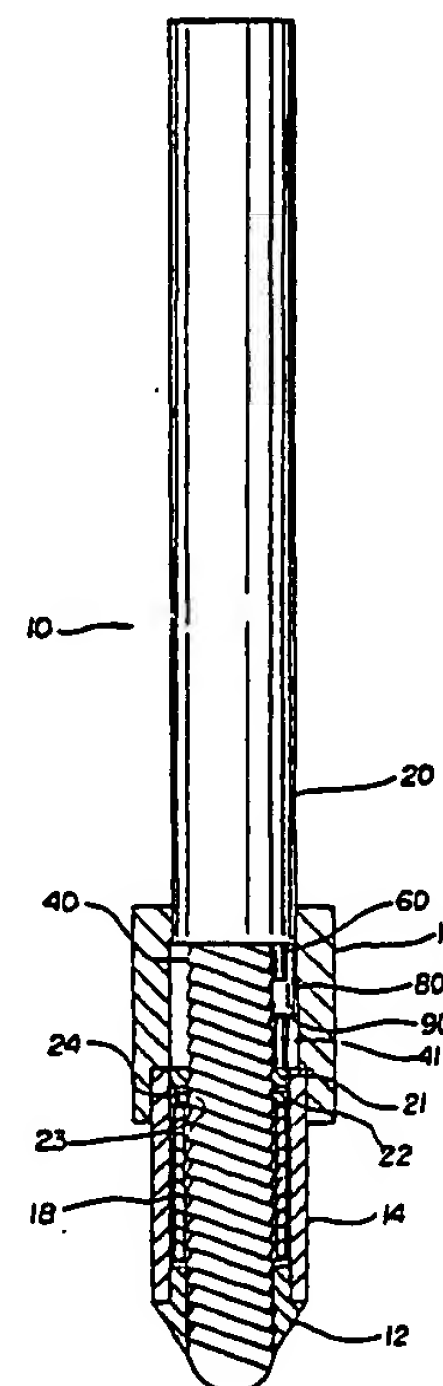
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61F 2/06		A3	(11) International Publication Number: WO 96/39998
			(43) International Publication Date: 19 December 1996 (19.12.96)
(21) International Application Number: PCT/US96/05842		(81) Designated States: CA, JP, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(22) International Filing Date: 26 April 1996 (26.04.96)			
(30) Priority Data: 08/484,006 7 June 1995 (07.06.95) US		Published With international search report. With amended claims.	
(71) Applicant: SCIMED LIFE SYSTEMS INC. [US/US]; One Scimed Place, Maple Grove, MN 55311-1566 (US).		(88) Date of publication of the international search report: 23 January 1997 (23.01.97)	
(72) Inventors: DEL TORO, Connie; 2165 Shenandoah Court, Unit H, Plymouth, MN 55447 (US). PETERSON, Dean, A.; 2904 17th Avenue North, Minneapolis, MN 55411 (US). SHOEMAKER, Susan, M.; 11106 190th Avenue N.W., Elk River, MN 55330 (US).		Date of publication of the amended claims: 13 March 1997 (13.03.97)	
(74) Agents: ANDERSON, William, E., II et al.; Vidas Arrett & Steinkraus, Suite 1540, 920 Second Avenue South, Minneapolis, MN 55402-4014 (US).			

(54) Title: PULL BACK SLEEVE SYSTEM WITH COMPRESSION RESISTANT INNER SHAFT

(57) Abstract

A stent delivery system for delivering a stent comprising, a stent disposed on the distal end of the catheter, and an inner core, wherein the inner core is resistant to appreciable compression or accordion. The catheter further comprises a first sheath covering a portion of the inner core, wherein at least a portion of the distal end of the inner core is left uncovered by the first sheath, a retractable sheath which covers at least a portion of the stent and a portion of the distal end of the inner core and a retracting means for retracting the distal sheath to release the stent. The present invention also discloses a method of making the retractable sheath for a stent delivery system, comprising the steps of placing a sheath made of material comprising tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins on a mandrel, winding a wire coil around the Teflon® sheath, heating the tubing, cooling the tubing and removing the mandrel and coil.



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AMENDED CLAIMS

[received by the International Bureau on 11 February 1997 (11.02.97);
original claims 1-20 replaced by amended claims 1-19 (4 pages)]

1. A stent delivery system comprising:
 - a substantially incompressible inner core having proximal and distal ends, wherein the inner core comprises a spring coil;
 - 5 a stent concentrically arranged around the inner core near the distal end;
 - a retractable sheath surrounding at least a portion of the stent, and retracting means connected to the distal sheath for retracting the distal sheath;
 - 10 whereby when the retracting means is retracted the distal sheath is retracted freeing the stent for delivery.
2. The stent delivery system as in claim 1, further comprising an outer sheath having proximal and distal ends and covering a portion of the inner core, wherein at 15 least a portion of the distal end of the inner core is left exposed by the outer sheath, the retractable sheath overlapping a portion of the distal end of the outer sheath.
3. The stent delivery system as in claim 2, wherein the retracting means comprises a pull back wire attached to the retractable sheath, the pull back wire being 20 longitudinally moveable.
4. The stent delivery system as in claim 3, further comprising an annular ring, wherein the pull back wire is attached to the annular ring and wherein the retractable sheath is adhered to the annular ring allowing the pull back wire to retract the 25 retractable sheath.
5. The stent delivery system as in claim 4, further comprising a stopper positioned to prevent the stent from moving proximally as the retractable sheath is retracted to expose the stent.

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6. The stent delivery system as in claim 5, further comprising a pull back wire lumen which partially encloses the pull back wire and is at least partially covered by the outer sheath.
- 5 7. The stent delivery system as in claim 6, further comprising a stiffening wire having proximal and distal ends positioned longitudinally along the inner core.
8. The stent delivery system as in claim 7, wherein the distal end of the stiffening wire and the distal end of the pull back wire lumen are truncated prior to the annular
10 ring.
9. The stent delivery system as in claim 4, wherein the retractable sheath comprises a proximal sheath having proximal and distal ends, the proximal end of the proximal sheath overlapping the distal end of the outer sheath, and a distal sheath
15 having proximal and distal ends, the distal end of the proximal sheath being connected to the proximal end of the distal sheath and wherein the annular ring is attached to the proximal end of the distal sheath.
10. The stent delivery system as in claim 9, further comprising a manifold, the
20 manifold comprising a sliding member which is connected to the pull back wire and acts to retract said pull back wire.
11. The stent delivery system as in claim 10, wherein the outer sheath comprises polyolefinic ionomer material.
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12. The stent delivery system as in claim 11, wherein the retractable sheath is made of material which comprises tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins.
- 30 13. The stent delivery system as in claim 12, wherein the distal sheath comprises a contoured pattern of grooves.

14. The stent delivery system as in claim 13, further comprising a hydrating luer located on the manifold, a distal tip located at the distal end of the catheter and a communication means between the hydrating luer and the distal tip which allows for hydrating the distal tip from the hydrating luer.

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15. A retractable sheath for a stent delivery system comprising a tube of material made from tetrafluoroethylene fluorocarbon polymers or fluorinated ethylene-propylene resins and which is contoured with a pattern of grooves, the contoured sheath being made from the process comprising the steps of:

- 10 placing the tube on a mandrel, wherein the diameter of the mandrel is slightly smaller than the inside diameter of the tube;
- winding a wire coil around the tube;
- heating the tube;
- cooling the tube; and
- 15 removing the mandrel and coil.

16. A method of making a retractable sheath for a stent delivery system, comprising the steps of:

- placing a sheath made of material comprising tetrafluoroethylene
- 20 fluorocarbon polymers or fluorinated ethylene-propylene resins on a mandrel, wherein the diameter of the mandrel is slightly smaller than the inside diameter of the sheath;
- winding a wire coil around the sheath;
- heating the sheath;
- cooling the sheath; and
- 25 removing the mandrel and coil.

17. A method as in claim 16, wherein the wire coil is a cross-section wire coil.

18. A method as in claim 17, wherein the wire coil is a rectangular wire coil.

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19. A stent delivery system comprising:

a substantially incompressible inner core having proximal and distal ends, and being made from a wire coil with a predetermined pitch such that it is flexible in an uncompressed state, and will become rigid when compressed;

5 a stent concentrically arranged around the inner core near the distal end;

a retractable sheath surrounding at least a portion of the stent, and retracting means connected to the distal sheath for retracting the distal sheath;

10 whereby when the retracting means is retracted to free the stent for delivery, the distal sheath is retracted, causing the wire coil to compress and taking up the excess pitch in the wire coil, making the wire coil substantially rigid, preventing the inner core from collapsing.

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